In the claims

- (Currently Amended) A process for producing <u>polynucleotide phosphorylase</u>
 (PNPase), comprising at least the following steps:
- (A) constructing an expression vector comprising a prokaryote-derived polynucleotide phosphorylase (PNPase) gene, which gene is isolated from a prokaryote selected from the group consisting of Escherichia coli and its analogous bacteria and is integrated into a plasmid having a T7 promoter as an expression-regulating signal;
- (B) transforming Escherichia coli or its analogous bacteria having a T7 RNA polymerase gene using the expression vector;
- (C) allowing the resulting transformant to express the <u>polynucleotide</u>

 <u>phosphorylase</u> (PNPase) gene thereby accumulating <u>polynucleotide phosphorylase</u> (PNPase) in
 the bacteria, <u>and further continuing to allow expression until the bacteria is disrupted to release</u>
 the <u>polynucleotide phosphorylase</u> (PNPase) into the <u>supernatant outside of the bacteria</u>; and
- (D) recovering the bacteria having PNPase accumulated therein, and extracting and purifying the polynucleotide phosphorylase (PNPase) released in the supernatant.
 - (Cancelled)
- (Currently Amended) The process according to claim 1, wherein the plasmid has
 a tag gene capable of adding a tag to the <u>polynucleotide phosphorylase</u> (PNPase) to be produced.
- 4. (Previously Presented) The process according to claim 3, wherein the tag gene is a His tag gene, T7 tag gene, S tag gene, Nus tag gene, GST tag gene, DsbA tag gene, DsbC tag gene, CBD_{cex} tag gene, CBD_{cenA} tag gene, CBD_{clos} tag gene, Trx tag gene, HSV tag gene, or 3×FLAG tag gene.

- (Currently Amended) The process according to any one of claims 1, 3 and to 4, 11 or 12, wherein the prokaryote is Escherichia coli.
- (Currently Amended) The process according to claim 5, wherein the Escherichia coli is Escherichia coli K12, <u>Escherichia coli C600K</u> or <u>Escherichia coli C157</u>.
- (Previously Amended) The process according to claim 1, wherein the Escherichia coli having a T7 RNA polymerase gene is Escherichia coli BL21 [DE3], Escherichia coli BL21 [DE3] pLysS, Escherichia coli BLR [DE3], Escherichia coli Rosetta [DE3], or Escherichia coli B834 [DE3].
 - 8-10. (Previously Cancelled)
 - 11-13. (Cancelled).
- 14. (Previously Presented) The process according to claim 3, wherein the Escherichia coli having a T7 RNA polymerase gene is Escherichia coli BL21 [DE3], Escherichia coli BL21 [DE3] pLysS, Escherichia coli BLR [DE3], Escherichia coli Rosetta [DE3], or Escherichia coli B834 [DE3].
- 15. (Previously Presented) The process according to claim 4, wherein the Escherichia coli having a T7 RNA polymerase gene is Escherichia coli BL21 [DE3], Escherichia coli BL21 [DE3] pLysS, Escherichia coli BLR [DE3], Escherichia coli Rosetta [DE3], or Escherichia coli B834 [DE3].
- (Previously Presented) The process according to claim 5, wherein the Escherichia coli having a T7 RNA polymerase gene is Escherichia coli BL21 [DE3], Escherichia coli BL21 [DE3] pLysS, Escherichia coli BLR [DE3], Escherichia coli Rosetta [DE3], or Escherichia coli B834 [DE3].

- 17. (Previously Presented) The process according to claim 6, wherein the *Escherichia coli* having a T7 RNA polymerase gene is *Escherichia coli* BL21 [DE3], *Escherichia coli* BL21 [DE3] pLysS, *Escherichia coli* BLR [DE3], *Escherichia coli* Rosetta [DE3], or *Escherichia coli* B834 [DE3].
 - 18-19. (Cancelled).
- (New) The process according to any one of claims 1, 3 and 4 wherein the prokaryote is Salmonella typhtmurium.
- 21. (New) The process according to claim 20, wherein the Escherichia coli having a T7 RNA polymerase gene is Escherichia coli BL21 [DE3], Escherichia coli BL21 [DE3] pLysS, Escherichia coli BLR [DE3], Escherichia coli BLR [DE3